

IMPROVEMENTS IN, OR RELATING TO, A METHOD OF
DISPENSING AND/OR A DISPENSER

5 This invention relates to a method of
dispensing and/or a dispenser.

 Current dispensers of, in particular,
 inhalable material in aerosol form, require the
10 aerosol generating mechanism (such as a nebulizer) to
operate in an inefficient mode and this results in
different sized doses as the size of the dose depends
upon how the generating mechanism is used. This is
wasteful of the drug and the variability in dose size
15 can result in under or over dosing.

 It is therefore an object of the present
invention to provide a method of dispensing and/or a
dispenser which will obviate or minimise the
20 foregoing disadvantages in a simple yet effective
manner, or which will at least provide the public
with a useful choice.

 Accordingly, in one aspect the invention
25 consists in a method of dispensing inhalable material
comprising the steps of indicating to a patient that
a breath should be taken, detecting the state of
breathing, and causing or allowing said inhalable
material to be supplied to the mouth or nose of the
30 patient during a selected part of the period of
inspiration.

 In a further aspect, the invention consists
in a dispenser comprising a supply means to supply
35 inhalable material in inhalable form, said supply
means being in connection with atmosphere and a mouth

Referring to the drawings, a dispenser and/or a method of dispensing are provided as follows. The dispenser of Fig.1 comprises a supply means to supply inhalable material in inhalable form. This may be, in particular, any known type of aerosol generator or aerosol propellant such as a nebulizer 1. The aerosol may be processed by baffles to remove particles above a selected size and particles removed may be fed back into the nebulizer 1.

Referring to Figs. 1 and 2, a conduit 2 is provided from the nebulizer 1 to a mouth or nose piece 3 and the conduit 2 may lead from a cloud chamber or aerosol reservoir 4. Thus, particles produced by the nebulizer 1 may pass through the reservoir 4 along the conduit 2 to the mouth piece 3. A pair of branch conduits 5 and 6 may lead from the conduit 2, the conduit 5 allowing air to be inhaled and the conduit 6 allowing air to be exhaled. A conduit 7 extends from the chamber 4. Flow control means are provided in each conduit and thus, a valve 8 is provided in conduit 7, a valve 9 in conduit 5 and a valve 10 in conduit 6. Associated with each valve 8, 9 and 10 is a non-return valve 11, 12 and 13 directed to allow air flow in only the desired direction. The valves 8, 9 and 10 are desirably electrically operated comprising, for example, solenoid valves.

Detecting means are provided to detect air flow at least in the conduits 6 and 7. Thus, a detector 14 is provided in conduit 7 and a detector 15 in conduit 6. Each detector 14 and 15 preferably comprises a bobbin 17 which may be biased, for

inhalation is detected.

5 A power "on" pulse circuit 23 is provided
and when the dispenser is turned "on" a signal is
forwarded from power "on" pulse circuit 23 and signal
circuitry 24 indicates by visual or audio means that
a breath should be taken by the user. The valves 9
and 10 are closed and valve 8 is opened. This allows
10 inhalation of aerosol formed by the nebulizer 1 in
the reservoir 4 by the drawing of breath through the
inlet port 25. The state of breathing is detected
and, for example, when the start of inhalation is
detected by the detecting means comprising the flow
sensor 14, timer 26 begins and after a present delay
15 the valve 8 is closed. At this time, timer 27 is
started and valve 9 is opened about this time and
this, together with the non-return valve 12, allows
inhalation of air through the inlet 28. After timer
27 has run for its present time, valve 9 is closed
20 and the inhale signals at 24 are turned off. The
exhale signals at 35 are then turned on and valve 10
is opened which, together with the non-return valve
13, allows exhalation through the outlet port 29.
When the start of exhalation is detected by the flow
25 sensor 15, timer 30 is started and after this has run
for its present time, the valve 10 is closed and the
exhale signals from generator 35 are turned off. The
inhale signals from generator 24 are then turned on
and the valve 8 opens to restart the cycle when
30 inhalation occurs.

In the embodiment of Figs. 3 and 4, a
compressed air nebulizer is used, which is able to be
switched on and off and the intermittent nature of
35 the operation allows selection of the running period
to achieve a desired result. A conduit 2 is provided

5 a count of the number of breaths taken and this may be pre-settable so that, for example, an alarm sounds or the dispenser stops delivering aerosol after the required number of breaths have been taken. The counter may begin incrementing when the start of the inhalation is detected.

10 A power on pulse circuit is provided and when the dispenser is turned "on" a signal is forwarded from power on pulse circuit 23 and signal circuitry 35 indicates by visual or audio means that a breath should be taken by the user. This allows inhalation of aerosol formed by the nebulizer 1 in the reservoir 4 by the drawing of breath through the inlet port 25. When the start of inhalation is detected by the detecting means comprising the flow sensor 15, timer 30 begins. After 30 has run for its pre-set time the exhale signals at 24 are then turned on, which indicates that the user should exhale and signals at 35 are turned off. When the start of exhalation is detected by the lower sensor 14, timer 26 and timer 27 are started. After timer 27 has run for its present time the aerosol generator is turned on. After timer 26 has run for its pre-set time, the aerosol generator is turned off and signals at 24 are turned off and signals at 35 turned on. When inhalation occurs, the timer 30 is started to repeat cycle.

30 In use, the nebulizer 1 can be used to generate an aerosol of material such as insulin but which clearly could comprise other materials such as, for example, drugs for use in the treatment of asthmatic conditions. The drug used may dictate whether a compressed air or ultrasonic nebulizer is used.

CLAIMS

1. A method of dispensing inhalable material
5 characterised by comprising, in combination, the
steps of indicating to a patient that a breath should
be taken, detecting the state of breathing and
causing or allowing said inhalable material to be
supplied to the mouth or nose of the patient during a
10 selected part of the period of inspiration.
2. A method of dispensing inhalable material as
claimed in Claim 1, characterised in that said
15 selected part of the period of inspiration is towards
the beginning of the period of inspiration.
3. A method of dispensing inhalable material as
claimed in either Claim 1 or Claim 2, characterised
in that said method further includes the step of
20 indicating to said patient when to exhale.
4. A method of dispensing inhalable material as
claimed in any one of the preceding Claims,
characterised in that said method further includes
25 the step of indicating to said patient when to inhale.
5. A dispenser comprising a supply means (1) to
supply inhalable material in inhalable form, said
supply means being in connection with atmosphere and
30 a mouth piece (3) and characterised by further
including flow control means (8) to allow or
substantially prevent flow between said supply means
and said mouth piece and said supply means and said
atmosphere, signalling means (24) to indicate to the
35 user when to inhale, detection means (14 and 15) to
detect the commencement of inspiration and/or

5 12. A dispenser as claimed in Claim 6 and any claim dependent thereon, characterised that control means are provided to control said aerosol generator means at least some operations of said control means being initiated by said detection means.

10 13. A dispenser as claimed in Claim 12, characterised in that a branch conduit (6) is provided to allow breath to be exhaled.

15 14. A dispenser as claimed in Claim 11, characterised in that preferably non-return valves (11, 12, 13) allow breathing in a single direction only in said conduit and said branch conduits.

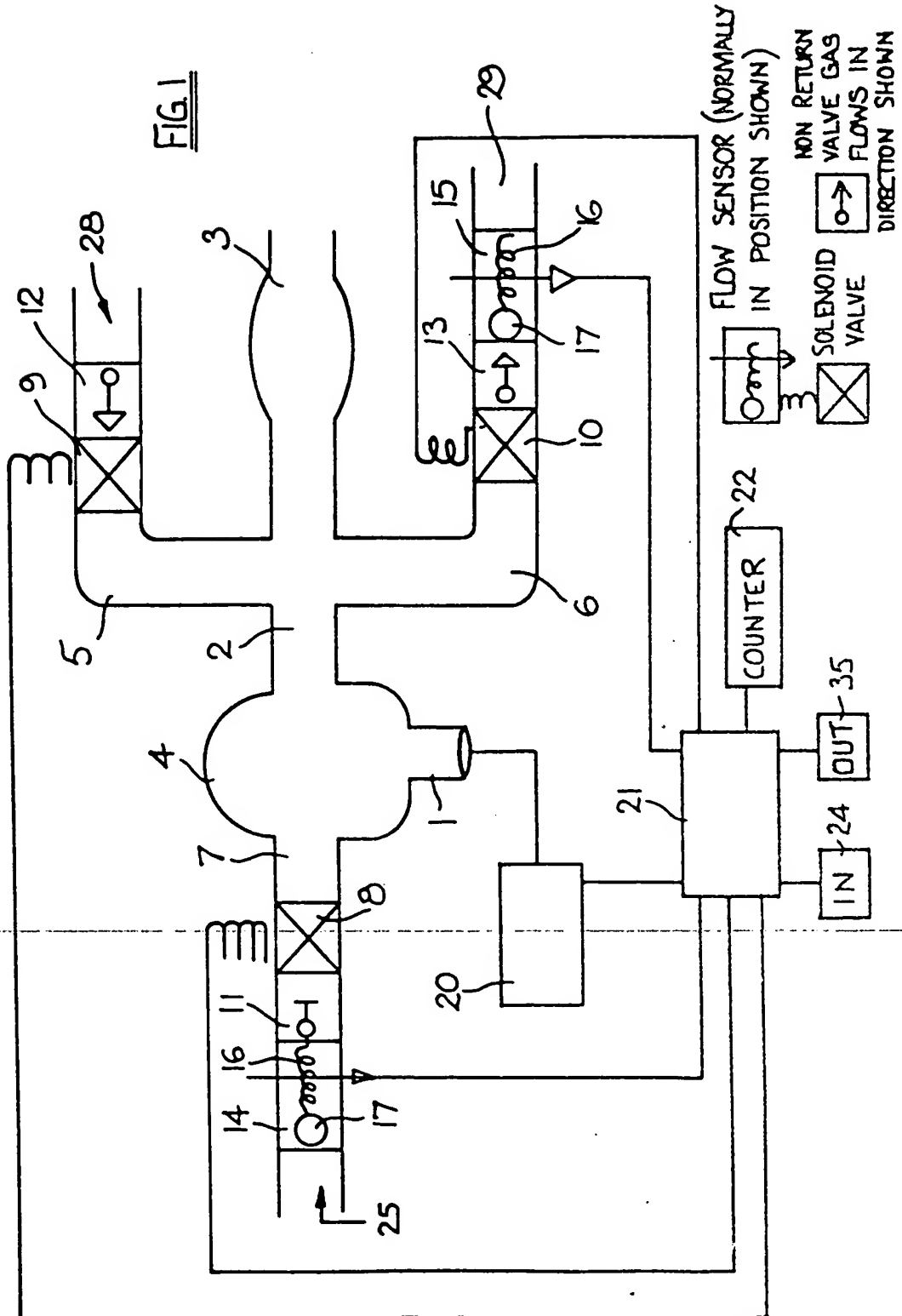
20 15. A dispenser as claimed in any one of Claims 5 to 14, characterised in that said detection means comprise a bobbin (17) moved by the breath of the patient and a light beam positioned so that said bobbin can interrupt said light beam.

25 16. A dispenser as claimed in any one of Claims 5 to 14, characterised in that said flow control means comprise one or more of a thermocouple or thermistor with a sensing circuit, a device to measure an increase or decrease pressure relative to atmospheric pressure, or a bobbin associated with a movement detector.

30 17. A dispenser as claimed in any one of Claims 5 to 16, characterised in that said signalling means (24, 35) comprise means to generate audible or visible signals to said user.

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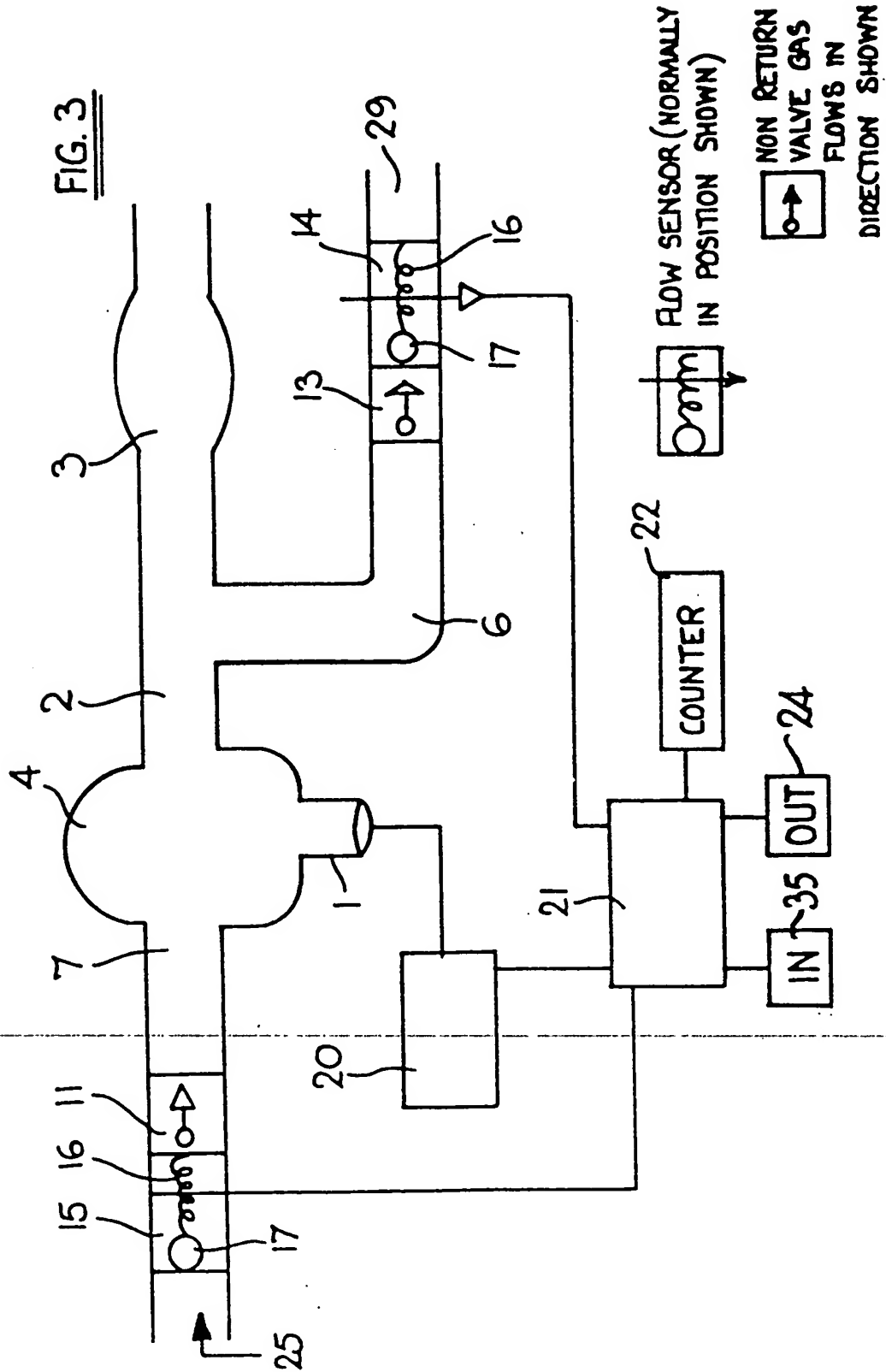
FIG. 1



[illegible]

OR GATE

FIG. 3



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